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Washington, DC 20231

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Sir:

Transmitted herewith for filing under 37 CFR 1.53(b) is:

☒ a patent application; **claiming the benefit of U.S. Provisional Application 60/080,383, filed 1 April 1998.**

☐ a Continuation    ☐ a Divisional    ☐ a Continuation-in-Part (CIP)

☒ A Small Entity Statement(s) was filed in the prior application; Status still proper and desired.

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Entitled: METHOD OF SUPPRESSING GENE EXPRESSION IN PLANTS

Enclosed are:

1. ☒ Application Transmittal Letter and Fee Transmittal Form (*A duplicate is enclosed for fee processing*)
2. ☒ 33 pages of Specification (including 35 claims)
3. ☒ 3 sheets of Informal Drawings (35 USC 113)
4. ☐ Oath or Declaration
  - a. ☐ newly executed (*original or copy*)
  - b. ☐ copy from prior application (37 CFR 1.63(d) (*for continuation/divisional*) [Note Box 5 Below]
  - c. ☐ **DELETION OF INVENTOR(S)** (*Signed statement deleting inventor(s) named in the prior application*)
5. ☒ Incorporation By Reference (*useable if box 4b is checked*)  
The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.
6. ☐ Microfiche Computer Program (*Appendix*)
7. ☐ Assignment papers (*cover sheet(s) and document(s)*) **Filed in parent application**
8. ☐ Small Entity Statement(s)
9. ☒ Information Disclosure Statement, PTO-1449, and 24 references cited
10. ☐ Preliminary Amendment (*Please enter all claim amendments prior to calculating the filing fee.*)
11. ☐ English Translation Document

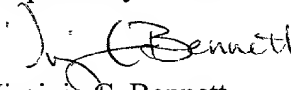
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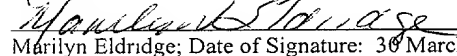
  
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## METHOD OF SUPPRESSING GENE EXPRESSION IN PLANTS

This application claims the benefit of U.S. Provisional Application No. 60/080,383, filed April 1, 1998.

### Field of the Invention

The present invention relates to the introduction of DNA episomes into plant cells to silence nuclear or chromosomal genes. More particularly, this invention relates to use of geminivirus vectors to provide  
5 systemic silencing of an endogenous plant gene in a treated plant.

### Background of the Invention

Gene silencing in plants typically refers to either the suppression of an endogenous gene by the introduction of a related transgene, or to the  
10 suppression of genes introduced at ectopic positions in plant genomes. Some examples of pathogen-derived host resistance to RNA viruses have been attributed to a gene silencing mechanism (Covey et al., *Nature* 386:781 (1997); Mueller et al. *Plant J.* 7:1001 (1995); Ratcliff et al., *Science* 276:1558 (1997); Tanzer et al., *Plant Cell* 9:1411 (1997). Transcriptional gene silencing has been hypothesized  
15 to involve DNA/DNA pairing, DNA methylation or heterochromatinization (Kumpatia et al., *Plant Physiol.* 115:361 (1997); Neuhuber, *Mol. Gen. Genetics* 247:264 (1995); Park et al., *Plant J.* 9:183 (1996)). Repeated DNA has a tendency to undergo transcriptional silencing, which may be associated with changes in chromatin structure (Meyer, *Biol. Chem. Hoppe Sayler* 377:87

(1996); Ye and Signer, *Proc. Natl. Acad. Sci. USA* 93:10881 (1996)), as well as to induce certain types of post-transcriptional silencing (Stam et al., *Plant J.* 12:63 (1997)). Both cytoplasmic and nuclear events have been implicated in gene silencing. Post-transcriptional gene silencing may require accumulation of a threshold level of mRNA, after which degradation of all homologous gene transcripts occurs, and may be potentiated by aberrant mRNAs. Metzlaff et al., *Cell* 88:845 (1997); Mueller et al., *Plant J.* 7:1001 (1995).

Geminiviruses are single-stranded DNA viruses that replicate through double-stranded DNA intermediates using the plant DNA replication machinery. Geminiviruses replicate in the nucleus, and foreign DNA can be stably integrated into the viral genome without significantly affecting replication or movement. Tomato golden mosaic virus (TGMV) is a bipartite geminivirus with a genome of two circular molecules, TGMV-A and TGMV-B (**Figure 1**). TGMV-A replicates autonomously. The TGMV coat protein (AR1) is dispensable for replication and movement in *N. Benthamiana* and can be replaced with up to 800 bp of foreign DNA, which is stably maintained in the viral genome (Elmer and Rogers, *Nucl. Acids. Res.* 18:2001 (1990)).

A plant virus may systemically infect a plant by spreading from the initially infected cell to neighboring cells, and subsequently throughout the plant. Plant cell walls prevent the random cell-to-cell transfer of virus, but channels (plasmodesmata) in plant cell walls provide an intercellular continuum through which virus particles or viral nucleic acids may move. Viral movement via plasmodesmata is mediated by virus-encoded proteins. Citovsky et al., *BioEssays* 13:373 (1991). Additionally, movement of virus to parts of the plant distant from the point of initial infection can occur via companion cells and sieve elements of the phloem. However, even in systemically infected plants the distribution of the virus may not be uniform. Certain areas of the plant, even within a tissue or a structure, may contain lower or higher amounts of virus than neighboring areas.

### **Summary of the Invention**

A first aspect of the present invention is a geminivirus silencing vector

comprising a geminivirus genome which contains heterologous DNA, where the heterologous DNA is identical to, or has substantial sequence similarity to, a gene endogenous to a plant or a fragment of a gene endogenous to a plant.

5 A further aspect of the present invention is a DNA construct comprising a geminivirus genome, where the DNA encoding the geminivirus coat protein has been replaced in part or in total with heterologous DNA identical to, or having substantial sequence similarity to, an endogenous plant gene or a fragment of an endogenous plant gene.

10 A further aspect of the present invention is a DNA construct comprising, in the 5' to 3' direction, a geminivirus origin of replication; DNA encoding proteins necessary for replication of the DNA construct; and a heterologous DNA segment identical to, or having substantial sequence similarity to, an endogenous plant gene or a fragment of an endogenous plant gene.

15 A further aspect of the present invention is a DNA construct comprising, in the 5' to 3' direction, a geminivirus origin of replication; DNA encoding proteins necessary for replication of the DNA construct; and a heterologous DNA segment identical to, or having substantial sequence similarity to, an endogenous plant gene or a fragment of an endogenous plant gene. This DNA construct is subject to the proviso that the DNA segment is not operably linked  
20 to a promoter.

A further aspect of the present invention is a method of silencing the expression of a plant gene in a plant cell, by inoculating the plant cell with a DNA construct or silencing vector as described above.

25 A further aspect of the present system is a method of systemically silencing expression of a plant gene in a plant, by inoculating the plant with an endogenous plant gene or a fragment of an endogenous plant gene.

30 A further aspect of the present invention is a method of screening isolated plant DNA for function. The method comprises preparing a DNA vector as describe above, containing a DNA segment identical to, or having substantial sequence similarity to, the isolated plant DNA or a fragment thereof. A test plant is then inoculated with the vector and allowed to grow for a period of time, then compared to a non-inoculated or sham inoculated control plant. Differences

between the inoculated and control plants indicate the function of the isolated plant DNA.

### **Brief Description of the Drawings**

**Figure 1A** shows the TGMV A and B genetic components; each contain a common region (CR) that includes the origin of replication. AL1, AL2 and AL3 are viral genes needed for replication and gene expression. The AR1 gene encodes the coat protein. The B component encodes two movement proteins, BL1 and BR1.

**Figure 1B** depicts the magnesium chelatase gene (*su*) cDNA, which includes 23 bp of upstream, non-coding sequence and a 1392 bp coding sequence. A 786 bp 5' fragment in sense orientation (*su*5S) was used to make TGMV::*su*5S. TGMV::*su*5F has a frameshift in the *su* coding sequence. TGMV::*su*5A contains a 786 bp 5' fragment in inverse (antisense) orientation. A 403 bp 3' fragment in sense orientation was used to make TGMV::*su*3S.

**Figure 1C** depicts the firefly luciferase gene (*luc*), along with two fragments derived from this gene used to make TGMV::*luc* constructs. A 623 bp fragment extending from the 5' end of the gene to the *Eco*R1 site was cloned into TGMV in sense (*luc*5S) and anti-sense (*luc*5A) orientations.

**Figure 2** shows the results of probing total RNA isolated from leaf and stem tissues to detect the accumulation of endogenous *su* mRNA. Lanes labeled "2 wk leaf" and "2 wk stem" show RNA harvested from plants 14 days post infection. "4 week" leaf tissue consists of leaves at a similar developmental stage as 2 week tissue (young, expanded), but taken from older plants (28 days post infection). Lanes 1, 5 and 10 contain RNA from uninfected control tissue; lanes 2, 6, and 9, TGMV::*su*5S-infected tissue; lanes 3 and 7, TGMV::*luc*5S-infected tissue; lane 4, wild-type TGMV-infected tissue; and lane 8, TGMV::*su*5F-infected tissue. TGMV::*su*-infected plants showed reduced accumulation of endogenous *su* mRNA. The episomal transcript is labeled TGMV::*su*5S and corresponds to the *su* transcript from the AR1 promoter of TGMV::*su*.

**Figure 3** shows the results of probing RNA harvested from plants 21 days post inoculation, with a *luc* RNA probe (*luc*), a probe for the constitutively

expressed *elF4A* gene (*elF4A*), and a probe for the constitutively expressed histone H1 gene (*histone*). Tissue was taken from two sham-inoculated plants (lanes 1-2), three plants infected with wildtype TGMV (lanes 3-5); three plants infected with TGMV::*luc5S* (lanes 6-8); and three plants infected with TGMV::*luc5A* (lanes 9-11). The luciferase transcript is downregulated by TGMV::*luc*. Degradation of the *luc* message in some of the samples is indicated by the smear below the full length *luc* transcript.

**Figure 4** shows the results of electrophoretic separation of uncut DNA probed with an 850 bp TGMV-A-specific DNA fragment. DNA was isolated from plants showing systemic symptoms or variegation; the tissue tested was either yellow (Y; yellow-white tissue); mixed (M; borders of green and yellow areas); or green (G; green tissue lacking visible traces of variegation or viral symptoms). Lane 1 shows wildtype TGMV DNA with three DNA forms: open circular (OC), closed circular (CC) and ssDNA. Lanes 2-4 show viral DNA isolated from *N. benthamiana* infected with TGMV::*su5S*; lanes 5-7 show viral DNA from plants inoculated with TGMV::*su5F*; lanes 8-10, viral DNA from plants inoculated with TGMV::*su5 A*. The bottom panel is the same blot as described above, but was probed with a 786 bp fragment from *su*. This gel indicates that TGMV::*su* DNA accumulation is lower in yellow tissue than in yellow-green borders.

#### **Detailed Description of the Invention**

The geminivirus tomato golden mosaic virus (TGMV) replicates in plant cell nuclei and expresses genes from high copy number DNA episomes. TGMV was used as vector to determine whether episomal DNA could cause silencing of homologous chromosomal genes. Two markers were used to assess silencing: (1) the sulfur allele (*su*) of magnesium chelatase, an enzyme required for chlorophyll formation; and (2) the firefly luciferase gene (*luc*). Various portions of both marker genes were inserted into TGMV in place of the coat protein open-reading frame and the constructs were introduced into intact plants using particle bombardment. When TGMV vectors carrying fragments of *su* (TGMV::*su*) were introduced into leaves of wild-type *Nicotiana benthamiana*, circular, yellow

spots with an area of several hundred cells formed after 3-5 days. Systemic movement of TGMV::su subsequently produced variegated leaf and stem tissue.

Fragments that caused silencing included a 786 bp 5' fragment of the 1392 bp *su* cDNA in sense and antisense orientation, and a 403 bp 3' fragment of *su* cDNA.

5 TGMV::su-induced silencing was propagated through tissue culture, along with the viral episome, but was not retained through meiosis. Systemic down-regulation of a constitutively expressed luciferase transgene in plants was achieved following infection with TGMV vectors carrying a 623 bp portion of *luc* in sense or anti-sense orientation.

10 The present results demonstrate that DNA carried on episomes can silence active, chromosomal gene expression, and that DNA plant virus genomes (and particularly geminivirus genomes) can provide a mechanism for the systemic suppression of gene expression in plants. The present inventors show that a nuclear-localized DNA virus (such as the TGMV geminivirus) carrying  
15 sequences complementary to (i.e., homologous to, or having substantial sequence similarity to) chromosomal genes can silence the chromosomal gene.

The present inventors determined that silencing of nuclear genes can be triggered by homologous sequences carried by a DNA episome, such as a geminivirus construct. Where the episome is capable of spreading from cell to  
20 cell in a plant (or capable of producing a diffusable silencing factor), systemic silencing of chromosomal genes can be achieved. A TGMV vector was used to test sense, anti-sense and frame-shifted versions of a 786 bp 5' fragment of the magnesium chelatase gene (*su*), as well as a non-overlapping 403 bp 3' sense fragment of *su*. In each case, similar variegated phenotypes were produced after  
25 infection. The lack of chloroplast transit sequences in the two *su* sense fragments used for silencing (the 5' sense fragment and the 3' fragment), as well as the anti-sense result, indicate that defective protein-protein interactions in the multisubunit magnesium chelatase complex are not responsible for the lack of chlorophyll formation. In addition, silencing of the luciferase transgene by  
30 TGMV::luc also appears to occur at the RNA or DNA level, as both sense and anti-sense versions of the *luc* fragment downregulate transgene expression. Luciferase was used to test downregulation by episomal sequences because its



activity can be visualized non-destructively. The activity of luciferase is known to be directly proportional to the accumulation of a single gene product (Millar et al, *Plant Cell* 4:1075 (1992)). Both luciferase activity and mRNA accumulation were markedly decreased in TGMV::luc-infected plants (see **Figure 4**).

5        Previous reports have demonstrated gene expression from geminivirus-derived episomes (reviewed in Timmermans et al., *Annu. Rev. Plant Physiol.* 45:79, 1994). A direct correlation between episome copy number and gene expression was shown in cultured cells for TGMV carrying the *neo* gene (Kanevski et al., *Plant J.* 2:457, 1992). The present experiments differ in that  
 10        there was no selection for gene expression, there was homology (i.e., sequence identity or substantial sequence similarity) between the episomal and chromosomal sequences, and only partial copies of the silenced endogenous genes (*su* or *luc* genes) were carried in episomes. The silencing of chromosomal gene expression by episomal DNA is demonstrated; more specifically, the ability  
 15        to silence endogenous gene expression systemically in a plant using a plant virus construct is demonstrated.

      The present invention permits the modulation of gene expression in plants, without the need for transformation of plants cells with subsequent regeneration of whole plants. The present methods also allow the modification  
 20        of plant gene expression without the modification of the germplasm.

#### Geminiviruses and Badnaviruses

      The geminiviruses are single-stranded plant DNA viruses. They possess a circular single-stranded (ss) DNA as their genome encapsidated in twinned  
 25        "geminata" icosahedral particles. The encapsidated ss DNAs are replicated through circular double stranded DNA intermediates in the nucleus of the host cell, presumably by a rolling circle mechanism. Viral DNA replication, which results in the simulation of both single and double stranded viral DNAs in large amounts, involves the expression of only a small number of viral proteins that are necessary  
 30        either for the replication process itself or facilitates replication or viral transcription. The geminiviruses therefore appear to rely primarily on the machinery of the host to copy their genomes and express their genes.

Geminiviruses are subdivided on the basis of host range in either monocots or dicots and whether the insect vector is a leaf hopper or a white fly species. Monocot-infecting geminiviruses are typically transmitted by leaf hoppers and their genome comprises a single ss DNA component about 2.7 kb in size (monopartite  
 5 geminivirus); this type of genome, the smallest known infectious DNA, is typified by wheat dwarf virus which is one of a number from the subgroup that have been cloned and sequenced. Most geminivirus that infect dicot hosts are transmitted by the white fly and possess a bipartite genome comprising similarly sized DNAs (usually termed A and B) as illustrated by African cassava mosaic virus (ACMV),  
 10 tomato golden mosaic virus (TGMV) and potato yellow mosaic virus. For successful infection of plants, both genomic components are required. Beet curly top virus occupies a unique intermediary position between the above two subgroups as it infects dicots but contains only a single genomic component equivalent to DNA A, possibly as a result of adaption to leaf hopper transmission.

15 The bipartite subgroup contains only the viruses that infect dicots. Exemplary is the African Cassava Mosaic Virus (ACMV) and the Tomato Golden Mosaic Virus (TGMV). TGMV, like ACMV, is composed of two circular DNA molecules of the same size, both of which are required for infectivity. Sequence analysis of the two genome components reveals six open reading frames (ORFs);  
 20 four of the ORFs are encoded by DNA A and two by DNA B. On both components, the ORFs diverge from a conserved 230 nucleotide intergenic region (common region) and are transcribed bidirectionally from double stranded replicative form DNA. The ORFs are named according to genome component and orientation relative to the common region (i.e., left versus right). The AL2 gene  
 25 product transactivates expression of the TGMV coat protein gene, which is also sometimes known as "AR1". Functions have not yet been attributed to some of the ORFs in the geminivirus genomes. However, it is known that certain proteins are involved in the replication of viral DNA (REP genes). *See, e.g., Elmer et al., Nucleic Acids Res.* 16:7043 (1988); Hatta and Francki, *Virology* 92:428 (1979).

30 The A genome component contains all viral information necessary for the replication and encapsidation of viral DNA, while the B component encodes functions required for movement of the virus through the infected plant. The DNA

5 information necessary for replication, encapsidation, and movement of the virus.

10 genome can trigger gene silencing in the host plant.

15 (ORF) may encode the movement protein, coat protein, protease and reverse transcriptase; the final products are produced by proteolytic processing.

### Episomally-mediated Gene Silencing

plant genes using DNA episomes, and provides constructs for the use in such methods. The episomal DNA carries a DNA sequence homologous to the endogenous plant gene to be silenced, or homologous to a fragment of the endogenous plant gene to be silenced. The DNA episomes must be able to replicate to multiple copy numbers in plant nuclei; where systemic silencing is desired, the episome should be able to move from cell-to-cell in the plant or to induce the movement of a diffusible suppression factor (or "silencing factor"), in order to enter and affect cells remote from the initial point of inoculation. The gene silencing may result in an altered phenotype; "altered phenotype" as used herein includes alterations in characteristics that can be visually observed (e.g., color), measured (e.g., average height or other growth characteristics) or biochemically assessed (e.g., presence of amounts of target gene products,

As used herein, an "endogenous" plant gene refers to a gene integrated into the chromosomal DNA of the plant gene. Endogenous genes include those that occur naturally in the plant genome, as well as those artificially introduced (such as by *Agrobacterium*-mediated transformation or ballistic bombardment).

As used herein, "systemic silencing" refers to the silencing of genes in plants or plant tissues, where gene silencing occurs in cells that are remote from the site of initial inoculation of the DNA silencing episome. Applicants do not wish to be held to a single theory of systemic silencing; systemic silencing may occur by the replication and cell-to-cell movement of DNA constructs, or by the movement of an as-yet-unidentified mobile silencing factor. Systemic silencing does not require that every tissue or every cell of the plant be affected, as the effects and extent of silencing may vary from tissue to tissue, or among cells.

- 10 -

upon introduction of the episome. Such DNA sequences are substantially similar in nucleotide sequence to the endogenous sequence to be silenced; the heterologous DNA sequence may have from 60% sequence similarity, 70% sequence similarity, 75% sequence similarity, 80% sequence similarity, 85% sequence similarity, 90% sequence similarity, 95% sequence similarity, or even 97% or 98% sequence similarity, or more, to the target endogenous sequence.

As used herein, the term "heterologous DNA" contained on the DNA silencing episome refers to DNA that is not naturally found in conjunction with the DNA episomal construct, i.e., that has been introduced by genetic engineering techniques. The heterologous DNA may be identical in sequence to an endogenous target gene to be silenced, or may have extensive sequence similarity thereto. The heterologous DNA may be identical in sequence to, or share extensive sequence similarity to, a fragment of an endogenous target gene, where the fragment is of a size sufficient to silence the endogenous target gene. The heterologous DNA may be in sense or antisense orientation. One skilled in the art will be able, using techniques available in the art and without undue experimentation, to test and select gene fragments for their ability to induce silencing when used in the present methods.

A preferred embodiment of the present invention utilizes DNA episomes based on plant viral genomes. A particularly preferred embodiment utilizes episomes based on geminivirus genomes. Additional plant DNA viruses include that Cauliflower mosaic virus and the Badnaviruses. Heterologous DNA may replace the coding region for the viral coat protein (or any other open reading frame that is nonessential for the present purposes of gene silencing).

Novel recombinant geminivirus constructs including expression cassettes and transfer vectors are provided. The present geminivirus constructs, when transfected into a plant cell, act to silence a gene already present in the plant cell. The gene to be silenced may be an endogenous plant gene, or a gene or DNA sequence that has previously been artificially introduced into the plant cell. The present geminivirus constructs further provide a method for the systemic silencing of a gene in a plant, for example, by providing both the A and B genome components of the geminivirus to the subject plant.

Also provided are methods of making the expression vectors and methods of using them to produce transfected plant cells having an altered phenotype. The expression vectors include a heterologous nucleic acid sequence as defined above. A preferred embodiment is a geminivirus genome DNA wherein the coding sequence of the coat protein gene has been replaced (in whole or in part) with the heterologous DNA sequence. Where a bipartite geminivirus genome is utilized, the DNA A genome component may include as operably linked components a transcription initiation unit (such as the geminivirus coat protein gene (AR1) promoter), a heterologous nucleic acid sequence as defined above, and a transcription termination region. The altered DNA A component may be transfected into a plant cell simultaneously with the DNA B component of that geminivirus; alternatively, the DNA-A and DNA-B components may be combined in a binary plasmid.

A transfected plant cell may be produced by contacting the plant cell with the recombinant DNA episome, such as a recombinant geminivirus transfer vector described above. Where a geminivirus construct containing both the DNA-A and DNA-B bipartite genome components is used, the components can be introduced into a plant cell on separate plasmids or carried together on a single DNA construct.

Thus the DNA constructs of the present invention include various forms, depending on the intended use of the construct. Constructs include vectors, transcriptional cassettes, expression cassettes, and binary plasmids. Where systemic silencing is desired, the construct must be capable of both replication in the host cell, and cell-to-cell movement (either of the DNA construct or a silencing factor).

A preferred recombinant episomal silencing construct contains a heterologous nucleotide sequence, which may be any sequence having sequence identity to, or substantial sequence similarity to, an open reading frame of an endogenous gene encoding a polypeptide of interest (for example, an enzyme).

Alternatively or in addition, the heterologous nucleotide sequence may be identical to or have substantial sequence similarity to an endogenous genomic sequence, where the genomic sequence may be an open reading frame, an intron, a

noncoding leader sequence, or any other sequence which inhibits transcription, messenger RNA processing, for example, splicing, or translation.

The heterologous DNA segment carried by the silencing construct may represent only a fragment of the endogenous gene to be silenced; the present  
5 inventors have shown that fragments of genes, in either the sense or antisense orientation, are sufficient to induce silencing.

A preferred DNA silencing construct comprises a DNA origin of replication, preferably from a plant DNA virus such as a geminivirus. The construct further includes DNA encoding any proteins necessary for replication of  
10 the DNA construct. For example, the origin of replication and DNA encoding necessary replication proteins may be obtained from the same geminivirus species; alternatively, the origin of replication may be from one geminivirus species and the replication proteins from a different geminivirus species. The construct further includes a DNA segment identical to, or having substantial sequence similarity to,  
15 an endogenous plant DNA segment to be silenced.

Episomal silencing constructs that utilize portions of DNA plant virus genomes do not have to include the viral movement protein genes to accomplish gene silencing. The present inventors have determined that the viral movement proteins are non-essential for episomally-mediated gene silencing. Because  
20 movement of the viral genome is not believed to be required for systemic silencing, the host plant need not be one that is naturally susceptible to the virus on which the silencing construct is based. Particle bombardment techniques, as are known in the art, may be used to introduce a silencing construct into a cell, or group of cells, in a plant. Replication of the construct in the plant cell(s) leads to  
25 systemic silencing. While not wishing to be held to a single theory, applicants believe that systemic silencing is achieved by movement of either the construct DNA, or a mobile silencing factor, through the plant tissues.

The nucleotide sequences of this invention may be synthetic, naturally derived, or combinations thereof. Methods of producing recombinant DNA  
30 constructs are well known in the art.

### Promoters

The present inventors have further determined that the DNA silencing episomes of the present invention need not have a promoter operably linked to the heterologous DNA segment therein. Use of a silencing construct carrying a heterologous DNA segment as described above, where that DNA segment is not operably linked to a promoter in the DNA construct, still results in silencing of the endogenous gene, and will result in systemic silencing. Use of a promoter operably linked to the heterologous DNA in the silencing construct may, however, increase the extent of the systemic silencing.

The present inventors have further determined that DNA constructs such as the geminivirus constructs described herein are capable of silencing endogenous plant promoters, where the DNA construct introduced into the target plant carries DNA having sequence identity to (or substantial sequence similarity to) an endogenous promoter sequence. Thus, where a family of homologous genes exist, but the associated promoters differ, selective silencing of one member of the gene family may be achieved by suppressing its promoter, using episomal constructs of the present invention. Such promoters may be tissue-specific (e.g., promoters associated with leaf-specific actins, as compared to actins expressed in other plant tissues) or developmentally regulated promoters. Examples of such promoters are known in the art.

### Methods

The present methods are useful in suppressing the production of undesired gene products (e.g., sugars or other products contributing to the flavor, color or composition of a plant product). Systemic or tissue-specific suppression of a particular endogenous protein product may be desirable where the plant has been transformed to express a protein product of interest; suppression of endogenous proteins may lead to increased production of the transgene of interest.

Improved methods of isolating and sequencing gene sequences have provided many isolated plant DNA segments of unknown function. Methods of determining the function of DNA segments have not kept pace with methods of isolating or determining the sequence of DNA segments. The present constructs



and methods provide a method of rapidly and reproducibly screening plant DNA fragments of unknown function to determine their function in plant tissue or intact plants, using episomally-mediated homology-dependent gene silencing. Such screening methods would include the preparation of an episomal silencing construct

5 containing heterologous DNA identical to or having substantial sequence similarity to the target sequence; inoculating host plants with the silencing construct and, after a period of growth, comparing the inoculated host with an uninfected control plant. Constructs based on geminivirus and badnavirus genomes are particularly useful, as these viruses are known to infect a wide variety of agriculturally

10 important crop plants. Characteristics for comparing test and control plants include growth characteristics, morphology, observable phenotype, and biochemical composition. The differences between the test and control plants indicate the function of the silenced DNA sequence. The period of growth necessary for any differences in the treated and control plants to become apparent

15 will vary depending on the host plants used and the function of the DNA being suppressed, as will be apparent to one skilled in the art. Such periods may range from several days, a week, two weeks, three weeks or four weeks, up to six weeks, eight weeks, three months, six months or more. Because the present method does not require tissue culture or selection to obtain alterations in gene

20 expression, the methods can be adapted to automation for large-scale screening of anonymous sequences for function in plants. As used herein, "screening" of a DNA segment does not imply that the function of the DNA segment will be positively identified in every case. As used herein, an "unidentified" plant gene or DNA segment is one whose functional role in the plant is unknown, even though

25 the nucleotide sequence may be known.

The present invention also provides methods for rapidly and reproducibly screening portions of an isolated plant gene of known function, to identify those portions or fragments of genes that are effective in preventing or suppressing expression. Such screening methods will lead to refinements in current methods

30 of gene suppression using sense and antisense DNA.

The present DNA episomal silencing system provides advantages over RNA viral vectors that are currently in use for testing gene function. Infection

with RNA viruses requires that infectious transcripts be made *in vitro*, capped, and mechanically inoculated. Other "knock-out" systems in plants rely on chromosomal transformation, which can be time-consuming. Unlike RNA virus-derived vectors, foreign DNA is stably maintained in geminivirus vectors and cloned DNA isolated from *E. coli* can be used directly for inoculation of intact plants, e.g., by particle bombardment. Infectious DNAs can be easily generated from shuttle vector libraries containing large segments of cDNA sequence. The present inventors have shown that as little as 150 base pairs of transcribed sequence can result in effective silencing, obviating the need for cloning full-length cDNAs. Promoter sequences have also been silenced by TGMV vectors, indicating that individual members of gene families can be selectively silenced where their promoters differ sufficiently from one another. Geminivirus and badnavirus vectors can be developed for different families of plants, thus allowing genes to be characterized directly in a species of interest. The present invention can also be used to identify single gene traits in a variety of species. Libraries of genes can be tested by subjecting plants to a screen for a single gene trait, such as pathogen resistance, and then looking for susceptible plants whose gene for resistance has been silenced. Current mutagenesis techniques require screening of segregating progeny, which can be time-consuming and is not feasible for many species that carry genes of interest.

#### Subject Plants

Plants which may be employed in practicing the present invention include any plant (angiosperm or gymnosperm; monocot or dicot) in which DNA constructs according to the present invention can replicate and, where systemic silencing is desired, where movement of the DNA construct or a silencing factor occurs. Particularly preferred are those plants susceptible to infection by plant geminiviruses. As used herein, "susceptible to infection" includes plants that are naturally infected by geminiviruses in the wild, plants that can be mechanically inoculated with the DNA construct, or that can be inoculated by methods other than mechanical inoculation (such as by *Agrobacterium* inoculation). "Susceptible

to infection" refers to plants in which the DNA construct is able to replicate within the inoculated plant cell.

Suitable plants for use in the present methods include (but are not limited to) legumes and solanaceous species. Specific plants suitable for use in the present invention included, but are not limited to, tobacco (*Nicotiana tabacum*), 5 potato (*Solanum tuberosum*), soybean (*glycine max*), peanuts (*Arachis hypogaea*), cotton (*Gossypium hirsutum*), squash, sweet potato (*Ipomoea batatas*), cassava (*Manihot esculenta*), beets, sugar beets (*Beta vulgaris*), and tomato (*Lycopersicon esculentum*).

A variety of techniques are available in the art for introduction of DNA constructs into a plant cell host. These include *Agrobacterium*-mediated transfection, injection, electroporation, microparticle bombardment, etc. In a preferred embodiment, intact plants are inoculated using microprojectiles carrying a geminivirus construct according to the present invention, that includes both the A and B DNA components. The site of inoculation will be apparent to one skilled in the art; leaf tissue is one example of a suitable site of inoculation. Inclusion of the B component of the geminivirus genome results in systemic effects; gene silencing is seen at sites remote from the original point of inoculation.

The examples which follow are set forth to illustrate the present  
20 invention, and are not to be construed as limiting thereof.

### EXAMPLE 1

## Materials and Methods: Vector Construction

**Figure 1A** depicts the TGMV A and B genetic components; each contains a common region (CR) that includes the origin of replication. AL1, AL2, and AL3 are viral genes needed for replication and gene expression. The AR1 gene encodes the coat protein and was replaced with the heterologous DNA fragments shown in **Figures 1B** and **1C** to produce the TGMV-derived episomes used herein. The B component encodes two movement proteins, BL1 and BR1.

30 TGMV-A-derived vectors were constructed using the pMON1655 plasmid, a pUC-based plasmid with 1.5 tandem copies of TGMV-A containing the AR1 coding sequence replaced by a short polylinker, and retaining the AR1

promoter and terminator sequences. pl-VN44 is a full length 1392 bp cDNA of the nucleotide-binding subunit of magnesium chelatase isolated from *Nicotiana tabacum* cv. SR1 (Nguyen, *Transposon tagging and isolation of the sulfur gene in tobacco*, Ph.D. Thesis, North Carolina State University (1995)). Magnesium chelatase is a multi-subunit protein that catalyzes the insertion of magnesium into protoporphyrin IX (Jensen et al., *Molec. Gene Genetics* 250:283 (1996)). In tobacco, a mutated allele (*Su*) of one subunit causes the phenotype known as 'sulfur'. *Nicotiana tabacum* plants homozygous for this allele are yellow (*Su/Su*), and heterozygous plants are yellow-green (*Su/su*).

A 786 bp *Acc661/EcoRV* fragment from the 5' end of the *su* cDNA (including the start codon and 17 bp of 5' non-coding sequence) was cloned into pMON1655 in both orientations, resulting in two TGMV constructs: the 5' *su* fragment driven by the TGMV AR1 promoter in the sense orientation (TGMV::*su5S*) and the anti-sense orientation (TGMV::*su5A*).

The sense reading frame of the *su* gene was disrupted by cleaving and filling an *EcoNI* restriction enzyme site to create a stop codon 104 bp from the initiator ATG to produce TGMV::*su5F*.

TGMV::*su3S* contains a 403 bp *EcoRV* fragment in sense orientation that ends 108 bp upstream of the putative stop codon.

The plasmids TGMV::*luc5S* and TGMV::*luc5 A* contain a 623 bp *EcoR1/BglIII* fragment of the 5' end of the full-length (1660bp) firefly luciferase gene (Ow et al., *Science* 234:866 (1996)) cloned into pMON1655 in the sense and anti-sense orientations, respectively. This fragment of the firefly luciferase gene contains 26 bp of 5' non-coding sequence and retains the luciferase start codon.

A plasmid containing 1.4 tandem copies of the TGMV B component (pTG1.4B, Fontes et al., *Plant Cell* 8:405 (1994)) was used with the TGMV-A constructs for all inoculations described herein, unless otherwise noted.

Table 1 describes the TGMV-A constructs used for the experiments described herein.

TABLE 1

TGMV-A constructs	
Name	Heterologous insert
TGMV::su5S	786 bp fragment of 5' end of <i>su</i> gene, sense orientation
TGMV::su3S	403 bp fragment of 3' end of <i>su</i> gene, sense orientation
TGMV::su5A	786 bp fragment of 5' end of <i>su</i> gene, antisense orientation
TGMV::su5F	frameshifted <i>su</i> gene, sense orientation
TGMV::luc5S	623 bp fragment of 5' end of firefly luciferase gene, sense orientation
TGMV::luc5A	623 bp fragment of 5' end of firefly luciferase gene, antisense orientation

### Example 2

#### 5        Materials and Methods: Plant Infections, DNA and RNA Analysis

Three-week-old *N. benthamiana* plants were used for bombardment. The BIOLISTIC® Particle Delivery System (Bio-Rad, Hercules, California, USA) was used to infect *N. benthamiana* plants as described previously (Nagar et al., *Plant Cell* 7:705 (1995)). Total DNA from infected and healthy plants was  
10 isolated (Dellaporta et al., *Plant Molec. Biol. Rep.* 1:19 (1983)), blotted and probed with labeled DNA corresponding to the TGMV-A AL1, AL2, AL3 reading frames, or 786 bp of *su*. Total RNA was isolated from infected or mock-infected leaf and stem tissue as described (Kaufman et al., *Plant Physiol.* 78:388 (1985)). Glyoxylated RNA was fractionated, blotted and hybridized with  
15 RNA probes as described (Frances et al., *Plant Cell* 4:1619 (1992)).

Transgenic plants were imaged using a photon-counting imaging system (Hamamatsu Photonic Systems, Bridgewater, New Jersey, USA) as described (Millar et al., *Plant Cell* 4:1075 (1992)). Systemically infected or control leaves were excised and immediately submerged in a 0.01 mg ml<sup>-1</sup> p-luciferin solution  
20 for precisely 5 minutes before being placed in the imaging chamber.

### Example 3

#### Plants Infected with TGMV::su display a Variegated Phenotype

To determine if TGMV could be used as a vector to silence plant gene expression, we tested fragments of *su* inserted into TGMV-A in place of the AR1 coding sequence, transcribed from the AR1 promoter (Example 1 and Table 1, above). Sequences from *su* were used because down-regulation of the endogenous gene was likely to produce a visible phenotype (lack of chlorophyll).

Although the cDNA used was from *N. tabacum*, the sulfur alleles of *N. tabacum* and *N. benthamiana* showed 95% nucleotide sequence conservation over the 5' and 3' fragments used in this study (unpublished observations).

*N. benthamiana* plants were infected using particle gun bombardment to deliver cloned viral DNA. Wild-type TGMV produced chlorotic, irregular spots on inoculated leaves approximately 5 days after bombardment; leaf curling and chlorosis were apparent at 5-7 days. In contrast, leaves inoculated with TGMV::su5S showed discrete, round, yellow spots after 3-5 days (data not shown). Not all yellow spots produced by TGMV::su5S were circular, but most had a distinct border between green and yellow tissue that was absent in plants infected with wildtype TGMV and in TGMV vectors carrying *luc* fragments. 'Spots' produced by inoculation of TGMV::luc and wt TGMV occurred later, were infrequent, and had mixtures of green and yellow cells (data not shown). Individual spots produced by TGMV::su5S varied in size from 1 to 3 mm and had an area of cells lacking chlorophyll that corresponded to at least 500-800 epidermal cells.

Variegation was produced by systemic infection with the TGMV::su constructs. As new leaves emerged subsequent to inoculation, large yellow-white or green sectors developed, often emanating from vascular tissue (data not shown). The patterns of variegation differed from plant to plant and from leaf to leaf.

In many plants, yellow or white tissue was confined to areas adjacent to the veins. In other plants, large sectors of mesophyll tissue were affected. White tissue likely resulted from bleaching of carotenoids. Microscopic examination of fixed tissue showed no evidence of necrosis or cell death in areas

lacking chlorophyll. A cross-section of leaves from TGMV::su5S infected plants and control plants showed that TGMV::su5S-infected leaf had a uniform lack of chlorophyll, compared to a mock-inoculated control leaf. When both leaves were viewed with fluorescence to detect DAPI-stained DNA, both leaves

5 indicated the presence of intact nuclei.

Replacement of the AR1 gene by foreign DNA, such as *su* is known to attenuate symptoms (Gardiner et al., *EMBO J.* 7:899 (1988)). Wild-type TGMV also produced chlorotic lesions, but these lesions usually contained collapsed cells. A striking feature of the variegated plants was the stem tissue, which was

10 either striped yellow and green, uniformly yellow, or white (data not shown). Neither chlorotic lesions or any other visible symptoms (other than stunting) were seen on stems of wild-type TGMV-infected plants, although stem tissue of such plants occasionally contains collapsed cells (Nagar et al., *Plant Cell* 7:705 (1995)).

15 Systemic variegation required TGMV-B in addition to a TGMV-A construct. TGMV-B contains two genes required for viral movement. When TGMV::su5S was bombarded in the absence of the TGMV-B component, yellow spots were produced but systemic variegation was not detected (data not shown). Wild-type TGMV-A inoculated without TGMV-B did not show visible spots,

20 probably because the TGMV-B component is required for symptom formation (von Arnim and Stanley, *Virology* 196:288 (1992)).

#### Example 4

##### Variegation is Not Caused by Mutant Proteins

25 The mutant *Su* allele cloned from *N. tabacum* has a single mis-sense mutation causing an altered amino acid (Nguyen, *Transposon tagging and isolation of the sulfur gene in tobacco*, Ph.D. Thesis, North Carolina State University (1995)). This polypeptide has been hypothesized to stably interfere with the function of the heteromeric magnesium chelatase complex, causing a light green phenotype

30 in heterozygous plants. Translation of RNA from the wild type 785 bp *su* fragment included in TGMV:su5S may produce a truncated polypeptide that would similarly disrupt assembly of the heteromeric complex. To test this

hypothesis, we introduced a frameshift mutation into the *su* fragment 104 bp downstream of the start codon to produce TGMV::su5F. The wild-type *su* polypeptide is 424 amino acids, including the putative transit sequence. The frameshift mutation was predicted to yield a truncated polypeptide with only 35 amino acids homologous to the *su* protein. If translation were to re-initiate at the next AUG (position 99), it would produce a polypeptide lacking a transit sequence that would be unlikely to be imported into the chloroplast. There were no repeatable differences in the variegation produced in plants inoculated with TGMV::su5S or TGMV::su5F (data not shown) suggesting that defective protein interactions in the chloroplast were not responsible for the yellow phenotype.

Plants bombarded with TGMV::su5A or TGMV::su3S also produced variegation similar to that seen with TGMV::su5S. The 5' anti-sense and the 3' sense fragments lack sequences for functional transit peptides, yet each was as effective as the 5' sense fragment of *su* in producing yellow spots and systemic variegation. These results indicate that down-regulation of *su* gene function is more likely to result from interactions at either the RNA or DNA level, than from defective protein-protein interactions.

The original *su* cDNA was tested in a plasmid vector incapable of replicating in plant cells (pLVN44) to determine if TGMV was required for homology dependent silencing of *su*. No systemic variegation occurred and no yellow spots on the bombarded leaves were observed.

To determine if tissue recovered from TGMV::su-infected tissue retained the variegated phenotype, plants were regenerated from sectors of systemically infected *N. Benthamiana*. Explants were taken from yellow/white (no chlorophyll), variegated, or green tissue of plants infected with TGMV::su5S. Variegated plants were recovered from yellow/white or variegated tissue (results not shown). In addition, one plant regenerated from yellow/white tissue was fully green, with no indication of variegation. Unlike the variegated regenerants, the green plant lacked viral DNA, as determined by DNA gel blot hybridization analysis (data not shown).

To determine if the TGMV::su constructs had integrated into chromosomal DNA, regenerated plants that retained variegation were allowed to



set seed. Seed harvested from flowers with white stems and sepals (which are normally green) produced green, wild-type seedlings, as did all seed harvested from variegated plants. This result indicates that the TGMV::su sequences did not integrate into chromosomal DNA at significant frequencies and/or that any  
 5 chromosomal form of TGMV::su was not sufficient to cause silencing of the endogenous *su* alleles.

### Example 5

#### Endogenous Su mRNA Levels are Reduced in Yellow Sectors of TGMV/su-

#### 10 Infected Plants

Some phloem-limited geminivirus cause yellow mosaic leaf tissue. We considered the possibility that TGMV::su-induced variegation reflected altered symptomology of the engineered virus rather than a reduction of *su* mRNA.

TGMV::su-infected plants showed reduced accumulation of endogenous  
 15 *su* mRNA. Total RNA was isolated from particle bombarded leaf and stem tissue (control) and leaf and stem tissue from plants systemically infected with TGMV::su5S, TGMV::su5F; TGMV::luc, or wildtype (wt) TGMV. Leaf material from TGMV::su-infected plants contained some green tissue but the majority was yellow or white, while stem tissue was uniformly green (control,  
 20 wild-type) or white (TGMV::su5S and TGMV::su5F). As shown in **Figure 2**, 15µg of total RNA isolated from leaf or stem tissue was loaded in each lane. The blot was probed with a <sup>32</sup>P-labeled RNA transcript made from pLVN44 containing the *su* cDNA clone. The 18s rRNA band served as a loading control. The episomal transcript is labeled TGMV::su5S and corresponds to the *su*  
 25 transcript from the AR1 promoter of TGMV::su. Lanes labeled "2 wk leaf" and "2 wk stem" show RNA harvested from plants 14 days post infection. "4 week" leaf tissue consists of leaves at a similar developmental stage as 2 week tissue (young, expanded), but was taken from older plants (28 days post infection). Lanes 1, 5 and 10 show RNA from uninfected control tissue; lanes 2, 6, and 9,  
 30 TGMV::su5S-infected tissue; lanes 3 and 7, TGMV::luc5S-infected tissue; lane 4, wt TGMV-infected tissue; and lane 8, TGMV::su5F-infected tissue.

**Figure 2** shows that a transcript of the endogenous *su* gene at the

expected size was present in control and wt TGMV-infected stem tissue (lanes 1, 4, 5 and 10) but lacking in TGMV::su5S-infected leaf and stem tissue (lanes 2, 6 and 9). RNA from tissue infected with TGMV::su5S also contained a transcript corresponding to the viral 786 bp *su* fragment. Accumulation of this transcript was attenuated in leaf tissue at a later stage of viral infection (compare lanes 2 and 6 containing tissue two weeks' post-infection, to lane 9 containing tissue at 4 weeks' post-infection). Lane 8 shows that tissue infected with TGMV carrying the 786 bp fragment with a frameshift mutation also lacked the full length *su* transcript from the endogenous gene.

The level of endogenous *su* transcription was also tested in plants infected with TGMV::luc5S. TGMV::luc has no homology to plant DNA; it shows stable movement but attenuated symptoms. The level of *su* transcript in TGMV::luc-infected leaf material (**Figure 2**, lanes 3 and 7) was similar to that of the control (**Figure 2**, lanes 1, 5 and 10).

### Example 6

#### Downregulation of Transgene expression

We used transgenic luciferase plants to determine if episomal TGMV sequences could silence expression of a foreign gene. *N. benthamiana* plants stably transformed with *luc* driven by the 35S CaMV promoter were bombarded with TGMV::luc5S and TGMV::luc5A, containing a 623 bp 5' transcribed *luc* sequence in sense or anti-sense orientation. Leaves were imaged 14 days post inoculation and compared to leaves from *N. benthamiana* controls (inoculated with particles only) or those inoculated with wild-type TGMV. Photon density from low to high was displayed in photographic images as blue, green, yellow, red and white (not shown).

Areas with both high and low luciferase activity were found in leaves from uninfected control plants and in plants systemically infected with wt TGMV 2 weeks after inoculation (results not shown). Only low levels of luciferase activity were found in leaves infected with TGMV::luc5S and TGMV::luc5 A. The complete absence of high level luciferase activity was surprising as viral infections are non-uniform. These results indicate that luciferase transgene

expression is downregulated by TGMV::luc.

RNA from uninfected control and infected plants was analyzed to determine if the reduced luciferase activity was caused by a reduction in mRNA accumulation. **Figure 3** shows total RNA isolated from the same plants as described immediately above, 3 weeks after infection. RNA was isolated from the top three inches of each of the plants and included systemically infected leaves, stems and meristematic areas, but not inoculated leaves; 5 µg of total RNA loaded in each lane. Blotted RNA was probed with a <sup>32</sup>P-labeled *luc* RNA probe (see top panel of **Figure 3**) or probes for the constitutively expressed genes *elF4 A* (bottom panel, **Figure 3**) and histone H1 (bottom panel, **Figure 3**).

**Figure 3** shows RNA from two particle bombarded plants (controls; lanes 1-2); three plants each infected with wildtype (wt) TGMV (lanes 3-5); three plants each infected with TGMV::luc5S (lanes 6-8); and three plants each infected with TGMV::luc5A (lanes 9-11). Degradation of the *luc* message in some of the samples is seen in the smear below the full length transcript.

The amount of *luc* transcript was greatly reduced in plants bombarded with TGMV::luc compared to both control and wt TGMV-infected plants. Accumulation of the *luc* transcript in two of the wt TGMV-infected plants was higher than that of uninfected controls. Increased luciferase activity, detected by photon imaging, has also been noted in TGMV-infected plants 3-4 weeks after inoculation (unpublished observation), suggesting an enhancement of 35S CaMV promoter activity by TGMV.

To determine whether the reduction in *luc* transcript reflected a general reduction in cytoplasmic transcript accumulation, we examined the levels of mRNAs encoding elongation initiation factor 4A (*elF4A*) and histone H1. The *elF4A* gene is constitutively transcribed in meristematic, leaf, stem and root tissues of tobacco (Mandel et al., *Plant Mol. biol.* 29:995 (1996)). An RNA gel blot was hybridized with probes for *elF4A* from tobacco and histone H1 from pea (Gantt and Key, *Eur. J. Biochem.* 166:119 (1987)) (**Figure 3**). The level of both the *elF4A* and histone H1 transcripts were similar for wt TGMV-infected

and TGMV::luc-infected plant tissue. These results demonstrate that the TGMV::luc constructs cause a specific reduction in *luc* mRNA accumulation.

#### Example 7

##### 5      Tissue with Incomplete Silencing Contains Higher Viral DNA Levels

To determine whether a direct relationship existed between the presence of viral DNA and silencing, gel blot hybridization analysis of DNA isolated from leaves systemically infected with TGMV::su5S, TGMV::su5A or TGMV::su5F (Figure 4) was performed. Variegated leaf tissue was separated into three categories: green tissue (G), yellow tissue (Y) and mixed (M). Green tissue had no apparent silencing phenotype or viral symptoms. The yellow category included yellow and white tissue that lacked chlorophyll, while mixed tissue contained light green regions and borders between yellow and green tissue. Uncut DNA was electrophoresed, blotted to membranes and probed with an 850 bp TGMV-A-specific DNA fragment and results are shown in Figure 4. Lane 1 shows wildtype (wt) TGMV DNA with three DNA forms: open circular (OC), closed circular (CC) and ssDNA. Lanes 2-4 show DNA isolated from plants infected with TGMV::su5S. Lanes 5-7 show viral DNA from *N. benthamiana* inoculated with TGMV::su5F and lanes 8-10, TGMV::su5 A. The bottom panel of Figure 4 shows the same blot as above but probed with a 786 bp fragment from *su*.

Green, healthy tissue showed a low amount of viral DNA (Figure 4) indicating that some cells in the green tissue were infected. Viral DNA levels were higher in mixed tissues than they were in yellow tissue. The stability of the viral *su* insert was shown by a lack of smaller fragments corresponding to deleted forms of TGMV::su. Each of the TGMV::su constructs tested showed the same pattern, with the highest viral DNA levels in mixed tissues. This experiment was repeated several times with variable results. However, none of the plants ever had higher viral DNA levels in yellow tissue than in mixed tissue.

30      The ratio of ss DNA to ds DNA in plants infected with the TGMV::su constructs (which lack the AR1 coat protein gene) is lower than it is in wild-type TGMV infections (Figure 4). The AR1 protein encapsidates ssDNA, forming

5 In principle, viral DNA in the yellow tissue might have been degraded as  
a result of a host response to viral infection. However, microscopic examination  
of DAPI-stained vibratome sections from yellow-white sectors showed intact,  
healthy cells (results not shown) and viral DNA isolated from such tissue did not  
appear more degraded than DNA isolated from 'mixed' or green tissue (**Figure**  
10 **4**). Furthermore, ethidium bromide-stained gels of uncut total DNA from yellow  
tissue showed no signs of chromosomal DNA degradation (data not shown). To  
exclude wild-type contamination, the same blots were striped and reprobed with  
the 786 bp *su* fragment. This probe hybridized to both ssDNA and dsDNA from  
each of the TGMV::*su* constructs (**Figure 4**). These experiments suggest that *su*  
15 gene silencing is not simply proportional to viral DNA accumulation.

### Silencing Vector Based on TGMV

When the TGMV-A vector was bombarded into plants in  
30 conjunction with the TGMV-B component, silencing of the target plant genes was  
seen systemically in the plant (i.e., at sites remote from the site of bombardment).  
It is not yet clear whether the vector was moving through the plant, or whether a

diffusible component distinct from the vector was causing systemic silencing. Although the TGMV episome is believed to replicate in only a few plant cells, silencing of the target endogenous plant gene was extensive and occurred throughout vegetative and floral parts of the plant. Viral disease symptoms were  
5 greatly attenuated in plants infected with the present construct (compared to infection with a wild-type geminivirus).

The foregoing examples are illustrative of the present invention, and are not to be construed as limiting thereof. The invention is described by the  
10 following claims, with equivalents of the claims to be included therein.

THAT WHICH IS CLAIMED IS:

1. A geminivirus silencing vector comprising a geminivirus genome which contains heterologous DNA, said heterologous DNA having substantial sequence similarity to a gene endogenous to a plant.

2. A vector according to claim 1, wherein said heterologous DNA replaces a segment of the coding sequence for the geminivirus coat protein.

3. A vector according to claim 1, wherein said heterologous DNA is operably associated with a promoter.

4. A DNA construct according to claim 3, wherein said promoter is the promoter that is associated with said endogenous plant gene.

5. A DNA construct according to claim 3, wherein said promoter is the geminivirus coat protein promoter.

6. A vector according to claim 1, wherein said heterologous DNA is in sense orientation.

7. A vector according to claim 1, wherein said heterologous DNA is in antisense orientation.

8. A vector according to claim 1, wherein said heterologous DNA has substantial sequence similarity to a fragment of said endogenous plant gene.

9. A vector according to claim 1, wherein said heterologous DNA has substantial sequence similarity to the entire coding region of endogenous plant gene.

10. A vector according to claim 1, wherein said geminivirus genome is selected from the group consisting of Tomato Golden Mosaic Virus (TGMV) and African Cassava Mosaic Virus (ACMV).

11. The vector of claim 1, wherein expression of said heterologous DNA modifies an observable plant phenotypic trait.

12. A DNA construct comprising a geminivirus genome, wherein the DNA encoding the geminivirus coat protein has been replaced in part or in total with heterologous DNA having substantial sequence similarity to an endogenous plant gene.

13. A DNA construct according to claim 12, wherein said heterologous DNA is operably associated with a promoter.

14. A DNA construct according to claim 13, wherein said promoter is the promoter that is associated with said endogenous plant gene.

15. A DNA construct according to claim 13, wherein said promoter is the geminivirus coat protein promoter.

16. A DNA construct according to claim 12, wherein said heterologous DNA is in sense orientation.

17. A vector according to claim 12, wherein said heterologous DNA is in antisense orientation.

18. A vector according to claim 12, wherein said heterologous DNA has substantial sequence similarity to a fragment of said endogenous plant gene.

19. A vector according to claim 12, wherein said heterologous DNA has substantial sequence similarity to the entire coding region of endogenous plant gene.



20. A vector according to claim 12, wherein said geminivirus genome is selected from the group consisting of Tomato Golden Mosaic Virus (TGMV) and African Cassava Mosaic Virus (ACMV).

21. The vector of claim 12, wherein expression of said heterologous DNA modifies an observable plant phenotypic trait.

22. A DNA construct comprising, in the 5' to 3' direction, a geminivirus origin of replication; DNA encoding proteins necessary for replication of said DNA construct; and a heterologous DNA segment having substantial sequence similarity to an endogenous plant gene.

23. A DNA construct comprising, in the 5' to 3' direction, a geminivirus origin of replication; DNA encoding proteins necessary for replication of said DNA construct; and a heterologous DNA segment having substantial sequence similarity to an endogenous plant gene, subject to the proviso that said DNA  
5 segment is not operably linked to a promoter.

24. A DNA construct according to claim 22 or 23, further comprising at least one copy of a geminivirus DNA-B component.

25. A DNA construct according to claim 22 or 23, further comprising a transcription termination region located 3' to said heterologous DNA.

26. A DNA construct according to claim 22, further comprising a promoter operably linked to said DNA segment.

27. A DNA construct according to claim 26, wherein said promoter is the geminivirus AR1 promoter.

28. A DNA construct according to claim 26, wherein said promoter is the promoter that is associated with said endogenous plant gene.

29. A DNA construct according to claim 22 or 23, wherein said DNA segment is in sense orientation.

30. A DNA construct according to claim 22 or 23, wherein said DNA segment is in antisense orientation.

31. A plant cell comprising a geminivirus silencing vector according to claim 1, 12, 22, 23 or 24.

32. A plant comprising a plurality of plant cells according to claim 31.

33. A method of silencing the expression of a plant gene in a plant cell, comprising inoculating said plant cell with a vector according to claim 1, 12, 22, 23 or 24.

34. A method of systemically silencing expression of a plant gene in a plant, comprising inoculating said plant with a vector according to claim 1, 12, 22, 23 or 24.

35. A method of screening isolated plant DNA for function, comprising:  
preparing a vector according to claim 1, 12, 22, 23 or 24, wherein said heterologous DNA of the vector has substantial sequence similarity to said isolated DNA;

- 5       inoculating a plant with said vector;  
          comparing said inoculated plant to a control plant;  
wherein differences between the inoculated and control plant indicate the function of the isolated plant DNA.

## METHOD OF SUPPRESSING GENE EXPRESSION IN PLANTS

### Abstract

The introduction of DNA episomes into plant cells to reduce or prevent the expression of endogenous nuclear or chromosomal genes is described. Geminivirus vectors to provide systemic silencing of an endogenous plant gene in a treated plant are described.

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Fig. 1A

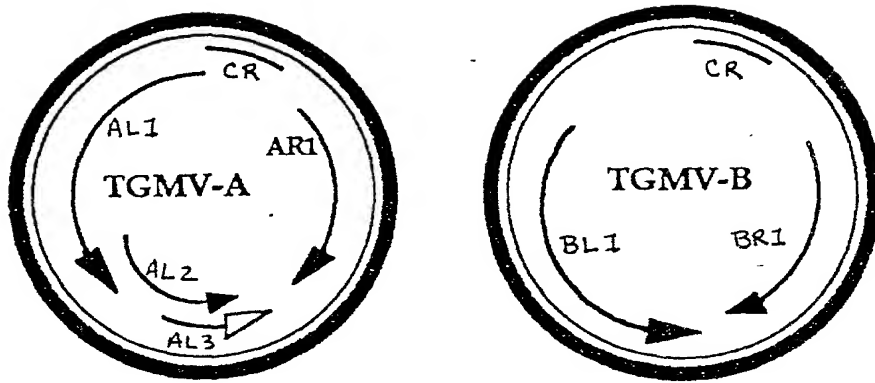


Fig. 1B

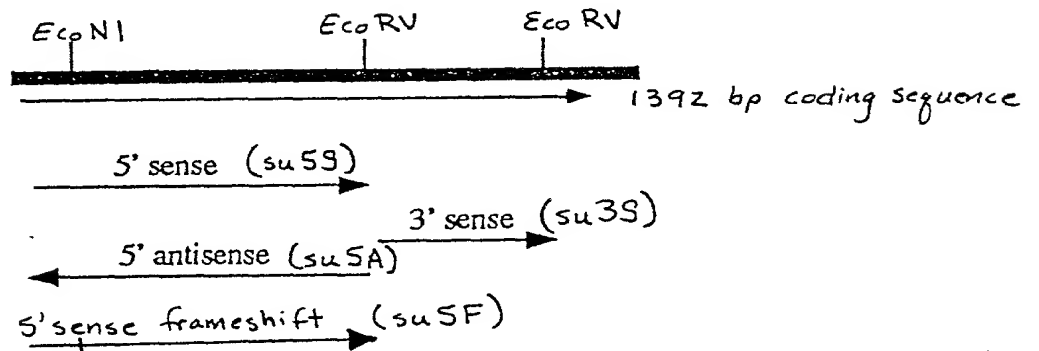
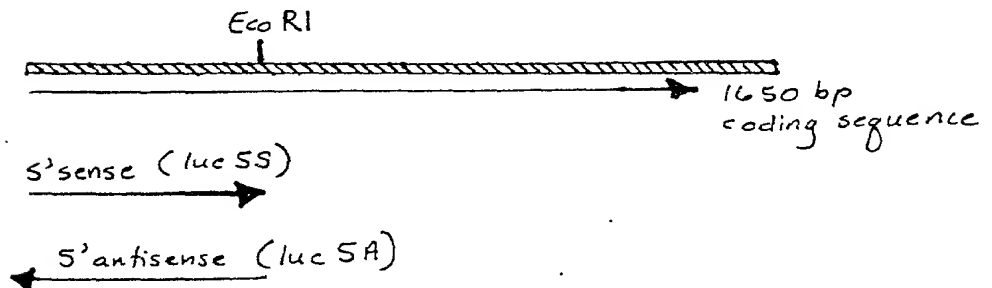


Fig. 1C



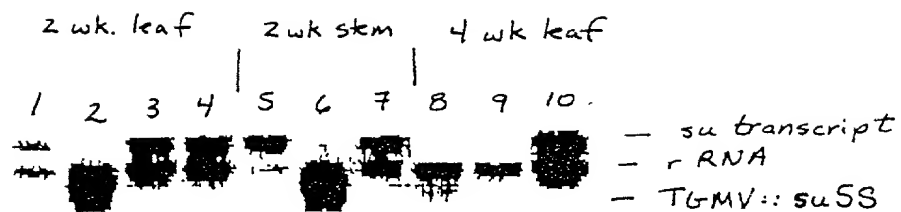


Figure 2

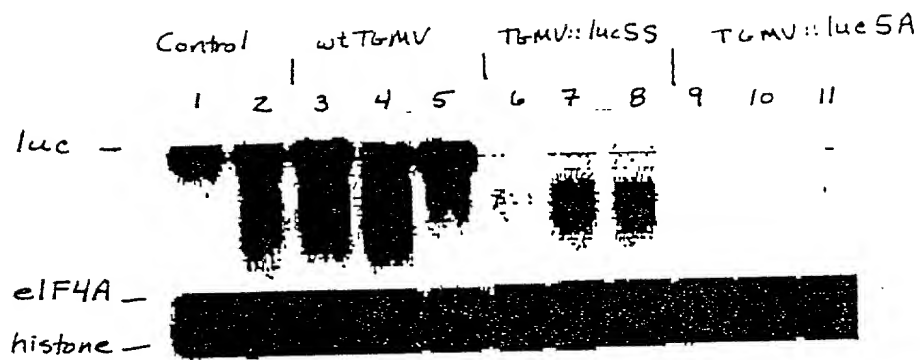


Figure 3

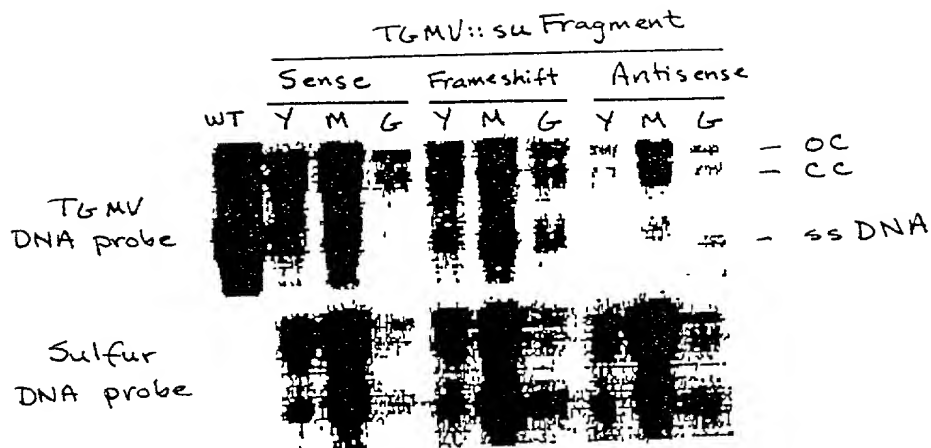


Figure 4